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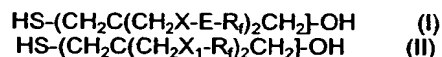
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(54) Heteroatom containing perfluoroalkyl terminated neopentyl mercapto-alcohols and compositions therefrom.

(57) Heteroatom containing perfluoroalkyl terminated neopentyl mercapto-alcohols of the formulas I and II



are prepared from the corresponding perfluorinated neopentyl oxetanes where R_f is a straight or branched chain perfluoroalkyl of 1 to 18 carbon atoms or said perfluoroalkyl substituted by perfluoroalkoxy of 2 to 6 carbon atoms, E is a linking group, and X and X_1 are heteroatom moieties.

The reaction products of these fluorinated mercapto-alcohols with electrophiles are disclosed as are certain functional derivatives. Compositions containing such materials provide improved thermal stability and useful low surface energy oil and water repellent coatings for textiles, glass, paper, leather, and other materials.

EP 0 501 915 A1

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Background of the Invention

This invention relates to novel hetero group containing perfluoroalkyl terminated neopentyl mercapto-alcohols which possess improved thermal stability. They are useful in the preparation of low surface energy oil and water repellent coatings for textiles, glass, paper, leather, and other materials.

One other bis-perfluoroalkyl terminated mercaptan has been previously described in U.S. Patent Nos. 3,758,447 and 3,886,201 and Japanese Sho 63-223,639. However, this compound is a perfluoroalkyl ester of mercaptosuccinic acid and is not only expensive to prepare, but also is not thermally and hydrolytically stable. Therefore, it is an impractical intermediate from which to obtain useful products.

The instant twin- R_f tailed mercapto-alcohols are readily isolated in high yield and purity. In addition, because of the neopentyl skeleton is incorporated into all of the novel mercapto-alcohols, they are thermally stable as well.

Bis-perfluoroalkyl mercapto-alcohols and derivatives thereof are useful because they possess a low free surface energy which provides oil and water repellency to a wide variety of substrates. Mercaptans containing a single R_f -function are known, but do not provide these properties to the same extent. The proximity of the two perfluoroalkyl chains to one another greatly increases the attraction between these groups through Van Der Waal's interactive forces, and as a result the oil and water repellency of the twin-tailed mercaptans is much greater than the repellency inherent in single-tailed mercaptans.

Objects of the Invention

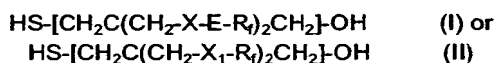
This invention most generally relates to novel heteroatom containing perfluoroalkyl terminated neopentyl mercapto- alcohols and compositions derived therefrom.

This invention also relates to the reaction products of said mercaptans with electrophiles. Other aspects of this invention are the derivatization of the alcohols as esters or urethanes. The products are useful per se or as intermediates for coatings on textiles, glass, linoleum, leather, wood, tile, metals, plastics and other various materials.

Another aspect of this invention relates to a substrate containing 0.01 to 50% by weight of a fluorine-containing mercapto-alcohol composition, at least part of said fluorine being provided by one unit derived from a heteroatom containing R_f -neopentyl mercapto-alcohol.

Detailed Disclosure

The instant invention pertains to novel heteroatom containing R_f -neopentyl mercapto-alcohols of formula I or II



wherein

R_f is a straight or branched chain perfluoroalkyl of 1 to 18 carbon atoms or said perfluoroalkyl substituted by perfluoroalkoxy of 2 to 6 carbon atoms,

E is branched or straight chain alkylene of 1 to 10 carbon atoms or said alkylene interrupted by one to three groups selected from the group consisting of -NR-, -O-, -S-, SO_2 -, -COO-, -OOC-, -CONR-, -NRCO-, $\text{SO}_2\text{NR-}$, and -NRSO₂-, or terminated at the R_f end with -CONR- or -SO₂NR-, where R_f is attached to the carbon or sulfur atom,

X is -S-, -O-, SO_2 -, or -NR-,

X_1 is -CONR- or -SO₂NR-, where R_f is attached to the carbon or sulfur atom, and

where

R is independently hydrogen, alkyl of 1 to 6 carbon atoms or hydroxyalkyl of 2 to 6 carbon atoms.

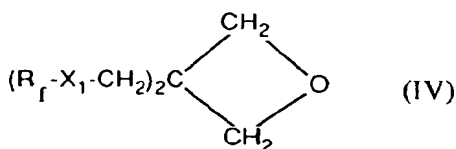
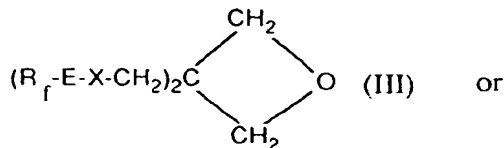
It is understood that the R_f group usually represents a mixture of perfluoroalkyl moieties. When the R_f group is identified as having a certain number of carbon atoms, the said R_f group also usually concomitantly contains a small fraction of perfluoroalkyl groups with a lower number of carbon atoms and a small fraction of perfluoroalkyl groups with a higher number of carbon atoms.

Preferably the instant compounds of formula I or II are those where R_f is perfluoroalkyl of 4 to 18 carbon atoms or perfluoroalkyl of 2 to 6 carbon atoms substituted by perfluoroalkoxy of 2 to 6 carbon atoms.

Preferably the instant compounds of formula I are those where R_f is perfluoroalkyl of 6 to 16 carbon atoms or perfluoroalkyl of 2 to 6 carbon atoms substituted by perfluoroalkoxy of 2 to 6 carbon atoms, E is alkylene of 2 to 6 carbon atoms, -CONHCH₂CH₂-, -CH₂CH₂N(CH₃)CH₂CH₂-, -CH₂CH₂SO₂NHCH₂CH₂-, -CH₂CH₂OCH₂CH₂-, or -SO₂NHCH₂CH₂-, and X is -S-, -SO₂- or -O-.

Most preferred are those compounds where R_f is perfluoroalkyl of 6 to 14 carbon atoms, E is ethylene, and X is S, i.e., $(R_fCH_2CH_2SCH_2)_2C(CH_2OH)(CH_2SH)$.

The novel R_f -mercapto-alcohols can be obtained directly by the reaction of a sulfur containing reagent with a bis- perfluoroalkyl oxetane of formula III or IV



The above-identified oxetanes are described in EP-A-0 430 887.

The synthesis of R_f -mercapto-alcohols proceeds by the nucleophilic addition of a sulfur containing reagent with the above-identified oxetane. This method of opening oxetanes to 3-mercapto-1-propanols has been previously described by Rondesvedt in the Journal of Organic Chemistry, 26, 3024 (1961). The reaction may be conducted in an aqueous system using phase transfer catalysis, but work-up of such an aqueous product is difficult due to troublesome emulsions. The improved process of this invention involves the combination of:

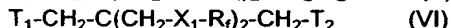
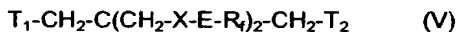
- a. an organic solvent, such as N-methylpyrrolidone, N,N-dimethylformamide, dimethyl sulfoxide, or the like, or alcohols such as methanol, ethanol, isopropanol or tert-butanol, or ketones, such as acetone, methyl alkyl ketones, or dialkyl ketones. Chlorinated solvents and esters generally give poor conversions and are less suitable;
- b. moderate reaction temperatures, on the order of 50 to about 120°C; and
- c. a stoichiometric quantity of a sulfur containing reagent, preferably an alkaline earth metal salt of thiocyanate, xanthate, mercaptoacetic acid or a neutral sulfur containing reagent such as thiourea, in the ratio of 1 mole of sulfur reagent per mole of oxetane to be cleaved; and
- d. a non-nucleophilic acid catalyst as exemplified by sulfuric acid or perchloric acid.

The reaction temperature and choice of solvent are mutually dependent. A reaction temperature in the range of 50-120°C is one wherein the formation of undesirable by-products is minimized and wherein the reaction products are stable.

Conditions are adjusted in order to achieve a reasonable rate of reaction at the chosen temperature.

Another aspect of the instant invention is a product prepared from the instant mercapto-alcohol by reaction with various electrophiles.

The heteroatom containing perfluoroalkyl terminated oxetanes described in EP-A-0 430 887 are not only precursors for the instant mercapto-alcohols, but also for the corresponding halohydrins and dihalides. These latter materials as well as the instant mercapto-alcohols can be used to prepare compounds of the formula V or VI



where

T_1 is the group VII



where

R_1 , R_2 , R_3 and R_4 are independently hydrogen, alkyl of 1 to 18 carbon atoms, hydroxyalkyl of 1 to 4 carbon atoms, haloalkyl of 1 to 4 carbon atoms, alkoxyalkyl of 2 to 12 carbon atoms, phenyl, phenyl substituted by one or two alkyl of 1 to 4 carbon atoms, by alkoxy of 1 to 4 carbon atoms, by carboxy or by hydroxy, phenylalkyl of 7 to 9 carbon atoms, carboxy, -COOZ where Z is alkyl of 1 to 18 carbon atoms, said alkyl interrupted by one or more -O- groups, allyl, cyclohexyl, phenyl, benzyl, tolyl or naphthyl; or -CONZ₁Z₂ where Z₁ and Z₂ are independently hydrogen, alkyl of 1 to 12 carbon atoms, hydroxyalkyl of 2 to 4 carbon atoms, phenyl, benzyl, alkoxyalkyl of 2 to 12 carbon atoms, cyclohexyl or Z₁ and Z₂ together are pentamethylene or 3-oxapentamethylene, with the proviso that at least one of R_1 to R_4 contains a carboxy, -COOZ or -CONZ₁Z₂ group, and

T_2 has the same meaning as T_1 or is hydroxyl.

Special interest is given to compounds of formula (V) or (VI) wherein R_f is a straight or branched chain perfluoroalkyl of 4 to 18 carbon atoms or perfluoroalkyl of 2 to 6 carbon atoms substituted by perfluoroalkoxy of 2 to 6 carbon atoms.

5 Preferred are compounds of formula V where

R_f is perfluoroalkyl of 6 to 16 carbon atoms or perfluoroalkyl of 2 to 6 carbon atoms substituted by perfluoroalkoxy of 2 to 6 carbon atoms,

E is alkylene of 2 to 6 carbon atoms, $-\text{CONHCH}_2\text{CH}_2-$, $-\text{CH}_2\text{CH}_2\text{N}(\text{CH}_3)\text{CH}_2\text{CH}_2-$, $-\text{CH}_2\text{CH}_2\text{SO}_2\text{NHCH}_2\text{CH}_2-$, $-\text{CH}_2\text{CH}_2\text{OCH}_2\text{CH}_2-$, or $-\text{SO}_2\text{NHCH}_2\text{CH}_2-$, and

10 X is $-\text{S}-$, SO_2- or $-\text{O}-$.

Most preferred are compounds of formula V wherein

R_f is perfluoroalkyl of 6 to 14 carbon atoms,

E is ethylene, and

X is $-\text{S}-$.

15 Preference is also given to compounds of formula (V) or (VI) wherein

R_f is a mixture of C_6F_{13} , C_8F_{17} , $\text{C}_{10}\text{F}_{21}$, $\text{C}_{12}\text{F}_{25}$ and $\text{C}_{14}\text{F}_{29}$.

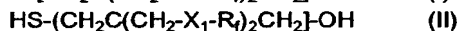
Preferably, one or both of T_1 and T_2 is a residue of a mercapto mono- or di-carboxylic acid of 2 to 4 carbon atoms after removal of the hydrogen atom from the mercapto group, and the other of T_1 and T_2 is hydroxyl.

20 Most preferably, both T_1 and T_2 are a residue of a mercapto mono- or di-carboxylic acid of 2 to 4 carbon atoms after removal of the hydrogen atom from the mercapto group; or T_1 is the residue of a mercapto mono- or di-carboxylic acid of 2 to 4 carbon atoms after removal of the hydrogen atom from the mercapto group, and T_2 is hydroxyl.

Preferably the mercapto acid is mercaptoacetic, thiolactic, 3-mercaptopropionic or mercaptosuccinic acid.

The compounds of formula V or VI where T_2 is hydroxyl and T_1 is a group VII can be made by

25 a) the addition of the mercapto-alcohol (I) or (II)



wherein X, X_1 , E and, R_f are defined as hereinbefore, to a compound of formula (X)

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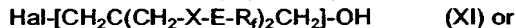


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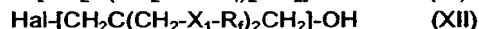
wherein R_1 , R_2 , R_3 and R_4 are defined as hereinbefore

or

b) by the reaction of an alcohol of the formula (XI) or (XII)



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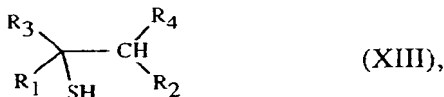
wherein

X, X_1 , E and, R_f are defined as hereinbefore and

Hal is halogen, preferably chlorine or bromine,

and a thiol of formula (XIII)

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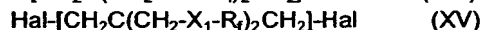
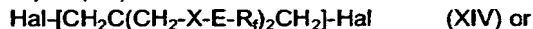
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wherein R_1 , R_2 , R_3 and R_4 are defined as hereinbefore.

The starting compounds of formula (XI) and (XII) are known from EP-A-0 430 887.

The compounds of formula V or VI where T_2 has the same meaning as T_1 can be made by the reaction of the bis-halogen compounds (XIV) or (XV)

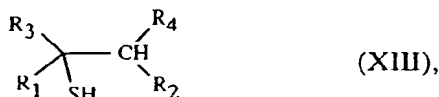
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wherein X, X_1 , E and, R_f are defined as hereinbefore, and

Hal is halogen, preferably chloro, bromo or iodo,

to a thiol of formula (XIII)



wherein R₁, R₂, R₃ and R₄ are defined as hereinbefore.

The starting compounds of formula (XIV) and (XV) are known from EP-A-0 430 887.

Some products can be derived through base-catalyzed nucleophilic additions to electrophilic substrates. For example, the base catalyzed addition reaction of mercaptans to alpha,beta-unsaturated esters is well known and described in detail in: Houben-Weyl, Methoden der Organischen Chemie, Volume 9/4, pages 123 to 126 (George Thieme Verlag, Stuttgart, 1955). Likewise, the base-catalyzed addition reaction of mercaptans to other electrophilic substrates, e.g. halides, sulfonates, acetates, is well known and described in Houben-Weyl, Methoden der Organischen Chemie, vol. 9/4 (Georg Thieme Verlag, Stuttgart, 1955, pp. 97-147), Barton and Ollis, Comprehensive Organic Chemistry vol. 3 (Pergamon Press, Oxford, 1979, pp. 13 - 16) and M. E. Peach in Patai, S., The Chemistry of the Thiol Group Part 2 (John Wiley and Sons, New York, 1974, pp. 722-734).

The types of electrophilic groups and hence the compounds that may be employed in conjunction with the perfluoroalkyl (R_f) containing mercapto-alcohols are numerous. The sole criticality is that nucleophilic addition/substitution occurs. Therefore in the presence of the disclosed mercaptans, addition/substitution will take place and the moiety will be introduced as a constituent of the final product. The basic reaction of mercaptans of the formula RSH involving addition/substitution of electrophiles is considered well documented in the literature.

Electrophiles of interest are generally all electrophiles which can be efficiently reacted and can therefore be employed for the synthesis of adducts. Several excellent surveys of mercaptan reactions with electrophiles are described and reference was made above to these teachings. A desired class of electrophiles therefore can be considered to be those possessing ethylenic unsaturation and those which readily undergo nucleophilic substitution.

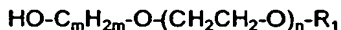
During the nucleophilic addition/substitution, the mercaptan derived moiety is introduced as an integral part of the molecule.

The class of unsaturated and other electrophilic compounds encompassed within the scope of this disclosure are those formed by compounds undergoing nucleophilic addition and contain the perfluoroalkyl moiety.

It is also possible to use free radical initiators or UV light for the addition of mercaptans to di- and triesters, as well as mono-esters and other nucleophilic substrates. This is possible because the di- and triesters are in contrast to acrylic esters, very reluctant to homopolymerize. A suitable catalyst may be any one of the commonly known agents for initiating the polymerization of vinyl monomers such as azo-initiators, (e.g., azobisisobutyronitrile) or aliphatic and aromatic acyl peroxides, e.g., decanoyl peroxide, lauroyl peroxide, benzoyl peroxide, dialkyl peroxides e.g., t-butyl peroxide, cumylperoxide; or hydroperoxides, e.g., t-butyl hydroperoxide, cumene hydroperoxide, or peresters and peroxy carbonates, e.g., t-butyl perbenzoate.

The addition of the mercaptans to the di- and triesters is usually carried out in a solvent in which the reactants and preferably also the adduct are soluble at the reaction temperature employed. Suitable solvents are aliphatic or aromatic hydrocarbons such as heptane, benzene, toluene, etc; chlorinated or fluorinated aliphatic or aromatic hydrocarbons such as methylene chloride, chloroform, methyl chloroform, carbon tetrachloride, trichloroethylene, perchloroethylene, FREON's such as 1,1,2-trifluoro- 1,2,2-trichloroethane, etc., chlorobenzene, benzotrifluoride or hexafluoroxylenes, ketones, esters and ethers such as acetone, methyl isobutyl ketone, ethyl acetate and higher homologs, dialkyl ethers, tetrahydrofuran, ethylene glycol monomethyl or monoethyl ether, ethylene glycol dimethyl or diethyl ether, and mixtures of these ketones, esters or ethers with water.

Non-ionic hydrophilic and hydrophobic activated olefins for such Michael reactions are known per se and many are commercially available, and include for example, but without limitation, alkyl acrylates and alkyl methacrylates, such as methyl methacrylate, ethyl methacrylate, isobutyl methacrylate, hexyl methacrylate, and n-octyl methacrylate. The esters of acrylic acids include methyl, ethyl, propyl, butyl, 2-hydroxyethyl, 3-hydroxypropyl, 2-hydroxypropyl or 2,3-hydroxypropyl esters; also ethoxylated and polyethoxylated hydroxyalkyl esters, such as esters of alcohols of the formula:



wherein

R₁ represents hydrogen or methyl,

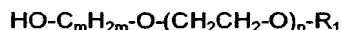
m represents 2 to 5 and

n represents 1 to 20, or esters of analogous alcohols, wherein a part of the ethyleneoxide units is replaced by propylene oxide units.

Further suitable esters are dialkylaminoalkyl acrylates and methacrylates, such as the 2-(dimethylamino)-ethyl-, 2-(diethylamino)-ethyl- and 3-(dimethylamino)-2-hydroxypropyl esters. The esters can be readily saponified with caustic to the parent acids if desired. Another class of hydrophilic electrophiles are acrylamide and methacrylamide as well as amides substitutes by lower hydroxyalkyl, lower oxaalkyl- or lower dialkylaminoalkyl groups such as N-(hydroxymethyl)-acrylamide and -methacrylamide, N-(3-hydroxypropyl)-acrylamide, N-(2-hydroxyethyl)-methacrylamide, N-(1,1-dimethyl-3-oxabutyl)-acrylamide and N-(1,1 -dimethyl-2-(hydroxymethyl)-3-oxabutyl)-acrylamide; methylol and ethers thereof: acrylamide, methacrylamide, diacetone acrylamide, and 2- hydroxyethyl methacrylate, as well as N-vinylpyrrolidone, acrylonitrile and methacrylonitrile.

Anionic hydrophilic activated olefins are known per se and include acrylic acid and methacrylic acid and salts thereof, acrylamidopropane sulfonic acid and salts thereof, maleic, fumaric, citraconic, muconic and itaconic acid and salts thereof as well as mono-olefinic sulfonic acids and their salts, such as sodium ethylene sulfonate, sodium styrene sulfonate and 2-acrylamido-2-methylpropane sulfonic acid.

Electrophiles which undergo nucleophilic substitution are, for example, but without limitation: halo-carboxylic acids and their derivatives such as chloro-, bromo- or iodoacetic, butyric, decanoic, hexadecanoic, hexanoic, dodecanoic, mandelic, octanoic, pyruvic, tetradecanoic, undecanoic, pivalic, valeric, succinic, and propionic acids. Ester derivatives include methyl, ethyl, propyl, butyl, 2-hydroxyethyl, 3-hydroxypropyl, 2-hydroxypropyl or 2,3-hydroxypropyl esters; also ethoxylated and polyethoxylated hydroxyalkyl esters, such as esters of alcohols of the formula:



wherein

R₁ represents hydrogen or methyl,

m represents 2 to 5 and

n represents 1 to 20 or esters of analogous alcohols, wherein a part of the ethylene oxide units is replaced by propylene oxide units.

Further suitable esters are dialkylaminoalkyl acrylates and methacrylates, such as the 2-(dimethylamino)-ethyl-, 2-(diethylamino)-ethyl- and 3-(dimethylamino)-2-hydroxypropyl esters. The above compounds can be substituted as well, for instance; chlorophenoxyacetic, 4-chloro-o-tolyloxyacetic, and 2-(2-chlorophenoxy)-2-methylpropionic acids. These esters can be readily saponified with caustic to the parent acids if desired.

Other anionic electrophiles are known per se and include 2-chloro-ethane sulfonic acid and salts thereof, 3-chloro-2-hydroxy-1-propanesulfonic acid and salts thereof.

If one of the components (reactant or solvent) boils below the reaction temperature, the reaction is desirably carried out in a pressure tube or an autoclave. It is most preferable for economic reasons to carry out the addition reactions in that solvent from which the adduct will be applied to a substrate such as textile, paper, leather and the like or which solvent can be utilized for additional reactions to be carried out with the adduct. The addition reaction is very simple to carry out, i.e. the mercaptan, the di- or triester are dissolved at the desired molar ratios in a solvent described above and the catalyst (0.01-2%) is added. The reaction mixture is kept at a temperature ranging from room temperature to 100°C, preferably under nitrogen until the disappearance of the double bond of the ester indicates that the reaction is complete. Other means of following the reaction are GC (gas chromatography), titration of free mercapto groups or TLC (thin layer chromatography). Required reaction times depend on reaction temperatures and amounts and kind of catalysis employed and may range from 5 minutes to 24 hours. To obtain products free of coloration, it is preferred to work at reaction temperatures below 80°C, preferably 40 to 70°C. If required, the addition product can be isolated by evaporating the solvent and catalyst (low volatile catalysts such as triethylamine are preferred) and be purified employing crystallization, precipitation or distillation procedures.

These mercapto-alcohols are characterized by the fact that they contain one pair of closely packed R_f-groups per SH-group. This fact is most important since closely packed R_f groups have been found to yield derivatives having higher oil repellency levels if compared with analogous derivatives derived from a mercaptan containing just one R_f-group or R_f-groups which are not closely packed, but separated by one or more backbone carbons.

The mercapto-alcohols can be used directly or indirectly with electrophiles to make a variety of condensation products and examples are presented in the experimental section.

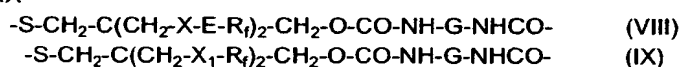
The novel adducts are useful in many ways: they can be applied to substrates such as textiles, paper, leather, wood, metallic surfaces and the like providing oil and water repellency to the treated substrates at extremely low add-ons. As shown in the examples, good repellency ratings are obtained with as little fluorine as 0.03 - 2% by weight of the substrate. The novel adducts may be applied to the various substrates by various coating techniques, such as dipping, spraying, brushing, padding, roll coating, and the like.

These adducts can be applied from a solvent and preferably from a solvent in which it was prepared for economical reasons. The adducts may also be applied from an aqueous system if either the adduct solution is water miscible or the adduct solution has been post-emulsified, employing emulsifiers and emulsification techniques known in the art.

Of course, it is understood that besides application to textiles, the coatings of the fluorinated compositions of the present invention are useful in providing oil and water repellent coatings for leather, paper, wood, masonry, metals, plastics, glass, painted surfaces, and the like. A very significant advantage of the compositions of the present invention is that they form effective oil repellent coatings at relatively very low fluorine levels on the substrate. In other words, on a given weight basis, the fluorine content of the compositions of this invention exhibits more effective repellent properties than the same level of fluorine in other compositions previously utilized in the art.

A further advantage of the compositions of this invention is that they may not require a separate curing or heating step, but can provide excellent repellent properties after drying in air.

Still another aspect of the instant invention are the polythiourethanes prepared from the instant mercapto-alcohols of formula I or II with a diisocyanate. Such polythiourethanes are represented by repeating structural units of formula VIII or IX



where

X, X₁, E, and R_f are as defined above in formula I and II and

G is the residue of an aliphatic, cycloaliphatic or aromatic diisocyanate after removal of the two NCO groups.

Preferred are polythiourethanes (VIII) or (IX) wherein

R_f is perfluoroalkyl of 4 to 18 carbon atoms or perfluoroalkyl of 2 to 6 carbon atoms substituted by perfluoroalkoxy of 2 to 6 carbon atoms.

Preferred are also polythiourethanes (VIII) wherein

R_f is perfluoroalkyl of 6 to 16 carbon atoms or perfluoroalkyl of 2 to 6 carbon atoms substituted by perfluoroalkoxy of 2 to 6 carbon atoms,

E is alkylene of 2 to 6 carbon atoms, -CONHCH₂CH₂-, -CH₂CH₂N(CH₃)CH₂CH₂-, -CH₂CH₂SO₂NHCH₂CH₂-, -CH₂CH₂OCH₂CH₂-, or -SO₂NHCH₂CH₂-, and

X is -S-, -SO₂- or -O-.

Mostly preferred are polythiourethanes (VIII) wherein

R_f is perfluoroalkyl of 6 to 14 carbon atoms,

E is ethylene, and

X is -S-.

Still another preferred embodiment are polythiourethanes (VIII) where

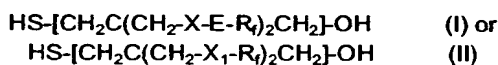
R_f is a mixture of C₆F₁₃, C₈F₁₇, C₁₀F₂₁, C₁₂F₂₅ and C₁₄F₂₉,

E is ethylene,

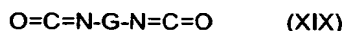
X is -S- and G is a radical of isophorone diisocyanate after removal of two NCO groups.

The polythiourethanes of formula VIII or IX can be prepared by reacting a mercapto-alcohol of formula I or

II



wherein R_f, X, E and X₁ are defined as before with an aliphatic, cycloaliphatic or aromatic diisocyanate of formula XIX



wherein

G is a bivalent aliphatic, cycloaliphatic or aromatic radical.

Any convenient diisocyanate can be used to react with the R_f-mercapto-alcohol. Myriads of useful diisocyanates are well-known in the art. Depending on the conditions selected an equivalent amount of a diisocyanate can be reacted with the mercapto-alcohol. Thus, one can use aliphatic, cycloaliphatic or aromatic diisocyanates.

Useful aromatic diisocyanates can be represented by the formula



where

G is the radical of the diisocyanate after removal of the two NCO groups.

Aromatic diisocyanates include:

tolylene diisocyanate (TDI) (all isomers), 4,4'-diphenylmethane diisocyanate (MDI), toluidine diisocyanate, dianisidine diisocyanate, m-xylylene diisocyanate, p-phenylene diisocyanate, m-phenylene diisocyanate, 1-

chloro-2,4-phenylene diisocyanate, 3,3'-dimethyl-4,4'-bisphenylene diisocyanate, 4,4'-bis(2-methylisocyanatophenyl)methane, 4,4'-bisphenylene diisocyanate, 4,4'-bis(2-methoxyisocyanatophenyl)methane, 1-nitro-phenyl-3,5-diisocyanate, 4,4'-diisocyanatodiphenyl ether, 3,3'-dichloro-4,4'-diisocyanatodiphenyl ether, 3,3'-dichloro-4,4'-diisocyanatodiphenylmethane, 4,4'-diisocyanatodibenzyl, 3,3'-dimethoxy-4,4'-diisocyanatodiphenyl, 2,2'-dimethyl-4,4'-diisocyanatodiphenyl, 2,2'-dichloro-5,5'-dimethoxy-4,4'-diisocyanatodiphenyl, 3,3'-dichloro-4,4'-diisocyanatodiphenyl, 1,2-naphthalene diisocyanate, 4-chloro-1,2-naphthalene diisocyanate, 4-methyl-1,2-naphthalene diisocyanate, 1,5-naphthalene diisocyanate, 1,6-naphthalene diisocyanate, 1,7-naphthalene diisocyanate, 1,8-naphthalene diisocyanate, 4-chloro-1,8-naphthalene diisocyanate, 2,3-naphthalene diisocyanate, 2,7-naphthalene diisocyanate, 1,8-dinitro-2,7-naphthalene diisocyanate, 1-methyl-2,4-naphthalene diisocyanate, 1-methyl-5,7-naphthalene diisocyanate, 6-methyl-1,3-naphthalene diisocyanate, 7-methyl-1,3-naphthalene diisocyanate, polymethylene polyphenyl isocyanate and co-products of hexamethylene diisocyanate and tolylene diisocyanate.

Useful aliphatic and cycloaliphatic diisocyanates include those of general formula



where G is the residue of the aliphatic or cycloaliphatic diisocyanate after removal of the two NCO groups.

Useful aliphatic or cycloaliphatic diisocyanates include:

1,2-ethane diisocyanate, 1,3-propane diisocyanate, 1,4-butane diisocyanate, 2-chloropropane-1,3-diisocyanate, pentamethylene diisocyanate, propylene-1,2-diisocyanate, 1,6-hexane diisocyanate, 1,8-octane diisocyanate, 1,10-decane diisocyanate, 1,12-dodecane diisocyanate, 1,16-hexadecane diisocyanate, 1,3-cyclohexane diisocyanate, 1,4-cyclohexane diisocyanate, 4,4'-methylene bis(cyclohexyl isocyanate) and isophorone diisocyanate.

Additionally, the following diisocyanates are particularly preferred because urethane compositions made therefrom tend to be non-yellowing:

1,6-hexamethylene diisocyanate (HDI), 2,2,4- and 2,4,4-trimethylhexamethylene diisocyanate (TMDI), dimer acid derived diisocyanate (DDI) obtained from dimerized fatty acids, such as linoleic acid, 4,4'-dicyclohexylmethane diisocyanate (hydrogenated MDI), isophorone diisocyanate, 3-isocyanatomethyl-3,5,5-trimethylcyclohexyl diisocyanate, lysine methyl ester diisocyanate (LDIM), bis(2-isocyanatoethyl) fumarate (FDI), bis(2-isocyanatoethyl)carbonate, m-tetramethylxylylene diisocyanate (TMXDI),

The reaction between the diisocyanate and the mercapto-alcohol component can be carried out in bulk, i.e., without solvent, or in the presence of non-reactive, anhydrous, organic solvents. Solvent media in which the reaction can be carried out include ketones, such as acetone, methyl ether ketone and methyl isobutyl ketone; esters such as ethyl acetate, isopropyl acetate, butyl acetate, 2-ethylhexyl acetate; hydrocarbons such as hexane, heptane, octane and higher homologs, cyclohexane, benzene, toluene, xylene or blends of aliphatic, cycloaliphatic and aromatic hydrocarbons or aprotic solvents such as N-methylpyrrolidine; it is also possible to employ ethers, both aliphatic and alicyclic including di-n-propyl ether, di-butyl ether, tetrahydrofuran and the diethers of polyalkylene oxides. In addition, chlorinated solvents such as 1,1,1-trichloroethane, dichloroethyl ether, ethylene dichloride, perchloroethylene and carbon tetrachloride can be used.

Among the solvents listed, the water miscible solvents such as acetone and methyl ethyl ketone are most important since they allow conversions of R_1 -urethanes into water soluble R_2 -urethanes as previously described.

In all cases, the solvents should be anhydrous to avoid urea formation.

The reaction can, if desired, be catalyzed and those catalysts conventionally employed in the urethane art are useful herein. Useful catalysts fall principally in two groups -

a. amino compounds and other bases:

triethylamine and other trialkylamines, triethylenediamine, 1,4-diaza-2,2,2-bicyclooctane, N-(lower)alkyl-morpholines, N,N,N',N'-tetra-methylethylenediamine, N,N,N',N'-tetramethyl-1,3-butanediamine, substituted piperazines, dialkylalkanolamines, benzyltrimethylammonium chloride and

b. organometallic and inorganic compounds:

cobalt naphthenate, stannous chloride, stannous octoate, stannous oleate, dimethyl tin dichloride, di-n-butyltin dilaurylmercaptide, tetra-n-butyl-tin, trimethyl-tin hydroxide, di-n-butyltin dilaurate.

Such catalysts may be used singly or in combination with each other. Beneficial synergistic catalysis may occur when combinations are used.

While it is possible to carry out the reaction without the use of a catalyst, it is preferable for reasons of economy and to assure a complete reaction, to utilize one or more catalysts as listed in amounts ranging from 0.01 to 1% based on the weight of the reactants. It is similarly advantageous to carry out the urethane synthesis at elevated temperature, usually between room temperature and 120°C, preferably at 60°C to 80°C, to obtain a complete reaction between 0.5 to 8 hours reaction time.

The reaction can be easily followed by titration of the isocyanate group or by IR analysis.

These end products include low molecular weight polythiourethane compositions useful to render plastics

soil repellent, and high molecular weight compositions useful as elastomers, foams, paints and varnishes, and textile treating compositions. The polythiourethanes are readily obtained by reaction of the said mercapto-alcohols with a variety of diisocyanates as described above.

These compositions have extremely low free surface energies and therefore, possess oil and water repellent properties, as well as mold release and other properties associated with low free surface energy. It should be noted that the compositions of this invention are characterized by the presence of two perfluoroalkylhetero groups in close proximity, a characteristic which provides improved oil and water repellent properties over the fluorinated compositions of the prior art. Further the two per-fluoroalkylthio groups are connected via a neopentyl moiety which does not permit the thermal elimination of mercaptan by beta-elimination. Hence, these R_f-mercapto-alcohols and derivatives have enhanced thermal stability.

Using the R_f-compounds and compositions described herein, it is possible to manufacture molds that display the excellent release properties characteristic of silicone oligomers. It is also possible to prepare polymeric urethane compositions with enhanced thermal stability.

The polythiourethanes are therefore useful as ingredients in floor polishes, furniture waxers, window washing fluids, and so on. Generally, they are useful as coatings on glass, ceramics, masonry, wood, plastics, textiles, leather and metals, or as additive ingredients in such coatings.

The usefulness of the polythiourethane compositions is, conveniently shown by measuring the oil, water and soil repellency ratings of substrates such as fabrics, paper, leather, etc. which are treated with solutions or emulsions of the novel polythiourethane compositions.

Coatings of the polythiourethane compositions may be applied to any desired substrate, porous or non-porous. They are particularly suited for application to porous materials such as textiles, leather, paper, wood, masonry, unglazed porcelain and the like to provide valuable oil and water repellency properties. However, they may also be applied to non-porous materials such as metals, plastics, glass, painted surfaces and the like to provide similar oil and water repellency properties. More specifically the polythiourethane compositions of the invention act as levelling, wetting and spreading agents in formulations designed for application to floors, furniture and automobiles. In such applications a protective oil and water repellent film is left on the treated object after the removal of the bulk of the material. Such levelling, wetting, spreading and film forming properties are also useful in

- a. formulations for cleaning glass and other hard, non-porous materials
- b. hair care products such as rinses, shampoos and hair sprays.
- c. paint, stain and varnish formulations for application to wood, masonry and ceramics.

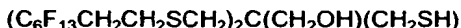
In the treatment of paper the polythiourethane compositions may be present as an ingredient in a wax, starch, casein, elastomer, or wet strength resin formulation. Aqueous emulsions of the polythiourethane compositions are especially useful in the treatment of paper. By mixing the polythiourethane compositions in an aqueous or oil type paint formulation, it may be applied effectively to unpainted siding, wood, metal and masonry. In the treatment of floors and tile surfaces and like substrates, the polythiourethane compositions may be applied by their incorporation in an emulsion or solution.

Because of the ability of the surfaces treated with these polythiourethane compositions to withstand abrasive action, the advantages incident to the repellency to oil and water and their resistance to soiling imparted by coating them with the polythiourethane compositions of this invention, preferred classes of articles to be treated are papers and textiles. Illustrative papers are carbonizing tissue, wallpaper, asphalt laminates, liner board, cardboard and papers derived from synthetic fibers.

Examples 1 to 6 and 8 to 22 illustrate the preparation of the R_f-mercapto-alcohols and derivatives. Examples 7, 25 and 26 illustrate the preparation of compositions, and certain utilities of these compositions. Examples 23 and 24 demonstrate the thermal superiority of the subject mercapto-alcohols.

The following examples are presented for the purpose of illustration only and are not to be construed to limit the nature or scope of the instant invention in any manner whatsoever.

Example 1



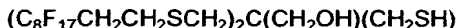
3,3-Bis(1,1,2,2-tetrahydroperfluorooctylthiomethyl)oxetane (100.0 gm, 0.12 mol) and thiourea (9.2 gm, 0.12 mol) are reacted under nitrogen with sulfuric acid (12.2 gm, 0.12 mol) and 2-butanone (125 gm) as the solvent. The reaction is carried out at 80°C for 4 hours. 2-(2-Aminoethylamino)ethanol (26.0 gm, 0.25 mol) is added over 0.5 hours. The product is stirred at 70°C and washed once with 350 ml distilled water. Residual water is removed as an azeotrope and the remaining solvent is evaporated under vacuum. The mercapto-alcohol (92 gm, 94% of theory) is of 89% purity by GLC. The crude product is recrystallized twice from 500 gm toluene to give a final product 95% pure by GLC, m.p. 63-65°C.

Analysis for $C_{21}H_{18}F_{26}OS_3$:

Calculated: C, 28.8; H, 2.1; F, 56.4; S, 11.0.

Found: C, 29.2; H, 2.0; F, 56.6; S, 10.7.

5 Example 2



3,3-Bis(1,1,2,2-tetrahydroperfluorodecylthiomethyl)oxetane (100.0 gm, 0.09 mol) and thiourea (7.3 gm, 0.10 mol) are reacted under nitrogen with sulfuric acid (9.6 gm, 0.10 mol) and 2-butanone (53.2 gm) as the solvent. The reaction is carried out at 80°C for 4 hours. 2-(2-Aminoethylamino)ethanol (26.0 gm, 0.25 mol) is added over 0.5 hours. The product is stirred at 70°C and washed once with 350 ml distilled water. Residual water is removed as an azeotrope and the remaining solvent is evaporated under vacuum. The mercapto-alcohol (95 gm, 94% of theory) is of 93% purity by GLC. The crude product is recrystallized twice from 500 gm toluene to give a final product 95% pure by GLC, m.p. 84-86°C.

15 Analysis for $C_{25}H_{18}F_{34}OS_3$:

Calculated: C, 27.9; H, 1.7; F, 60.0; S, 8.9.

Found: C, 27.5; H, 1.7; F, 58.6; S, 8.3.

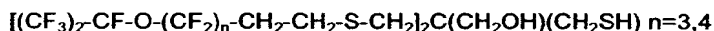
20 Example 3



N-(2-Mercaptoethyl)-perfluorooctanamide (11.3 gm, 0.025 mol) and 3,3-bis(bromomethyl)oxetane (3.1 gm, 0.013 mol) are reacted under nitrogen with potassium carbonate (3.3 gm, 0.025 mol) and 2-pentanone (4.0 gm) as the solvent. The reaction is run at 103°C for 19.5 hours. The reaction mixture is stirred and washed twice with distilled water. The solvent is evaporated to yield a yellow-brown wax. This product (15.0 gm, 0.016 mol) is then reacted with thiourea (1.3 gm, 0.017 mol), sulfuric acid (1.9 gm, 0.019 mol), and 2-propanol (18.2 gm) and heated to 90°C under nitrogen for five hours. The reaction is then reacted with 2-(2-aminoethylamino)ethanol (3.1 gm, 0.030 mol) and 2-propanol (19 gm) and heated to 85°C for fifteen minutes. The product is washed three times with distilled water and then collected. The mercapto-alcohol is of 85 % purity by GLC.

30

Example 4



1-Heptafluoroisopropoxy-1,1,2,2-tetrahydroperfluoro-alkanethiol (consisting of 73% n=3 homolog and 27% n=4 homolog) (3.7 gm, 0.009 mol) and 3,3-bis(bromomethyl)oxetane (1.1 gm, 0.005 mol) are reacted under nitrogen with sodium hydroxide (3mL of a 20% (w/w) aqueous solution). The reaction is carried out at 80°C for 3 hours. The product is a brown wax and is 87% pure by GLC. This material is then washed with water. The above oxetane is then reacted with thiourea (0.2 gm, 0.003 mol), sulfuric acid (0.3 gm, 0.003 mol), and 2-propanol (3.2 gm) as described in Example 1. The reaction is heated to 80°C for five hours under nitrogen. The product is a light-tan solid wax. This material is then reacted with 2-(2-aminoethylamino)ethanol (0.5 gm, 0.005 mol), and 2-propanol (3 gm). The reaction is heated for 15 minutes at 85°C. The product is a brown solid. The reaction is then washed with distilled water and the product is collected.

Analysis for product (n=3,4):

Calculated: C, 27.6; H, 1.9; F, 55.0; S, 10.3.

45 Found: C, 27.1; H, 1.4; F, 56.3; S, 9.3.

Example 5



50 4-Heptafluoroisopropoxy-1,1,2,2-tetrahydroperfluorobutanethiol (5.0 gm, 0.014 mol) and 3,3-bis(bromomethyl)oxetane (1.6 gm, 0.007 mol) are reacted under nitrogen with sodium hydroxide (4.7 ml of a 20% (w/w) aqueous solution). The reaction is carried out at 90°C for 3 hours. The product is a clear yellow oil which is 97% pure by GC. This material is then washed with water. The oxetane is then reacted with thiourea (0.3 gm, 0.004 mol), sulfuric acid (0.4 gm, 0.004 mol), and 2-propanol (3.4 gm) as described in Example 1. The reaction is heated to 90°C for five hours under nitrogen. The product is a brown solid wax. The isothiuronium salt is then treated with 2-(2-aminoethylamino)ethanol (0.73 gm, 0.007 mol) and 2-propanol (4 gm). The reaction is heated to 85°C for fifteen minutes. The product is then washed three times with distilled water and collected.

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Example 6Reaction Product of $R_fCH_2CH_2SCH_2)_2C(CH_2OH)(CH_2SH)$ and Maleic Acid

- 5 $(R_fCH_2CH_2SCH_2)_2C(CH_2OH)CH_2SCH(COOH)CH_2COOH$
 2,2-Bis-(1,1,2,2-tetrahydroperfluoroalkylthiomethyl)-propan-3-mercapto-1-ol* (1.0g, 0.9mmol), ditriethanolammonium maleate (0.4g, 0.9mmol), and Triton B (30 μ L of a 40% solution in methanol) are dissolved in 5mL of tert-butyl alcohol in a single neck 100mL round bottomed flask. The mixture is stirred at 50°C for 1.5 hour under nitrogen. The yellow solution which is formed dissolved easily in distilled water. The aqueous layer
 10 is acidified and extracted with methyl propyl ketone. The solvent is removed under vacuum and the resultant yellow oil is precipitated into water to produce a white solid. NMR shows proton resonances at 2.5ppm (4 x R_fCH_2); 2.6 - 3.0ppm (10 x SCH_2 ; 2 x CH_2CO_2); 3.51 (2 x CH_2OH); 3.72 (1 x $SCHCO_2$).
 Analysis for * R_f distribution C_6F_{13}/C_8F_{17} -9%/88%:
 Calculated: C, 29.0; H, 1.9; F, 53.3; S, 8.1.
 15 Found: C, 28.9; H, 1.7; F, 53.9; S, 8.2.

This material is formulated as the ditriethanolammonium salt in aqueous solution and evaluated for paper sizing as described below.

Sample Preparation and Testing

20

Pad Application of Fluorochemicals as an External Size

- A sample of the above fluorochemical is diluted to the test application levels with distilled water. The solution is added to a 4% aqueous solution of papermaker's starch and then applied to unsized paper by padding (paper
 25 dipped through starch solution, and passed through single nip rollers). The resulting sheets are dried on a hot metal surface (Williams sheet drier) 30 seconds both sides at 110°C.

Grease Resistance Test

- 30 Creased test papers are placed over a grid sheet imprinted with 100 squares. Five grams of sand are placed in the center of the crease. A mixture of synthetic oil and a dye for visualization is pipetted onto the sand and the samples are maintained at 60°C for 24 hours. Evaluation is determined by the percentage of the grid that stains.
 Another aspect of this invention describes the substrate containing 0.005 to 5% by weight of a fluorine-containing composition, at least part of said fluorine being provided by one or more units derived from the sub-
 35 ject R_f -chemicals.

Oil Resistance Test TAPPI UM 557

- 40 Samples of the fluorochemical product described in Example 6 are dissolved in water as a 20% (w/w) solution and are added to a 10% aqueous solution of papermaker's starch (pH adjusted to 9.6 with 10% aq. NaOH). This starch solution is applied to unsized paper by padding (paper dipped through starch solution, then passed through single nip rollers). The resulting sheets of paper are dried at ambient conditions for 15 minutes, then for 3 minutes at 200°C in an "Emerson Speed Drier" (heated metal plate with canvas cover).
 45 An easily made kit of twelve solutions of varying proportions of Castor Oil, Toluene, and Heptane is useful in comparing surface oil resistance.

50

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	<u>Kit No.</u>	<u>Volume Castor Oil</u>	<u>Volume Toluene</u>	<u>Volume Heptane</u>
5	1	200	0	0
	2	180	10	10
	3	160	20	20
10	4	140	30	30
	5	120	40	40
	6	100	50	50
	7	80	60	60
15	8	60	70	70
	9	40	80	80
	10	20	90	90
20	11	0	100	100
	12	0	90	110

The "kit value" is defined as the highest number solution that will stand on the surface of the plate for 15 seconds in the form of drops without failing. Failure is detected by pronounced darkening caused by penetration. The darkening of even a small fraction of the area under the drop is considered failure.

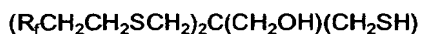
The AATCC Oil Rating is determined according to Standard Test method 118-1983 of the American Association of Textile Chemists and Colorists. Ratings are given from 0 (minimum) to 8 (maximum). A commonly accepted level of repellency for oil repellent fabrics in the United States is an oil repellency of 4.

Tests are run against the competitive commercial paper size - SCOTCHBAN FC-807 (3M Co.), a bis-perfluoroalkyl phosphate ester, ammonium salt. The products are applied to paper by pad application and tested for Oil Kit Rating and the Grease Resistance Test.

	<u>% Fluorine (Found).</u>	<u>Oil Kit</u>	<u>Grease Resistance</u>
<u>Test Compound</u>	<u>on Wt. of Paper</u>	<u>Number</u>	<u>Test</u>
Example 6	0.036	3	fail
	0.042	4	pass
SCOTCHBAN	0.060	6	fail
FC-807	0.066	7	pass

The results show that Example 6 has superior performance compared to that of the prior-art phosphate size, at much lower application levels, as measured by the Grease Resistance Test. Further, the novel instant carboxylate compound of Example 6 passes the Grease Resistance Test at lower Kit Numbers. This allows their application to products requiring better adhesive bonding, better label adhesion, and lessens problems with printing.

Example 7



3,3-Bis(1,1,2,2-tetrahydroperfluoroalkylthiomethyl)oxetane* (described in Example 1 of EP-A-0 430 887) is reacted with thiourea and sulfuric acid in 2-butanone as described in Example 1. The resultant isothiuronium salt is cleaved with 2-(2-aminoethylamino)ethanol to yield the desired mercapto-alcohol. The mercapto-alcohol is of 92% purity by GLC. m.p. 71-78°C.

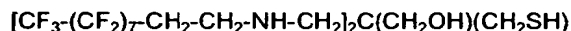
Calculated: C, 28.0; H, 1.7; F, 59.0; S, 9.0.

Found: C, 27.6; H, 1.7; F, 57.4; S, 8.2.

*R_F-distribution is C₆, C₈, C₁₀, C₁₂, C₁₄(%)-9/88/1.2/0.3/0.1

Example 8

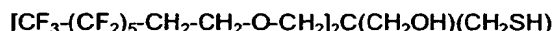
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3,3-Bis(bromomethyl)oxetane is reacted with 1,1,2,2-tetrahydroperfluorodecylamine in triethylamine as solvent at 80-100°C. The reaction is carried out for 12 hours and the product is worked up by acidification and precipitation into water. The powder is collected by filtration and dried. The oxetane is then treated with thiourea and sulfuric acid in tert-butanol and subsequently cleaved with 2-(2-aminoethylamino)ethanol as described in Example 1 to yield 2,2-bis-(1,1,2,2-tetrahydroperfluorodecyliminomethyl)-1,3-mercaptopropanol.

Example 9

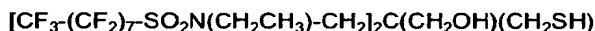
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3,3-Bis(1,1,2,2-tetrahydroperfluorooctyloxomethyl)oxetane (described in EP-A-0 430 887) is treated with thiourea and sulfuric acid in tert-butanol and subsequently cleaved with 2-(2-aminoethylamino)ethanol as described in Example 1 to yield 2,2-bis-(1,1,2,2-tetrahydroperfluorooctyloxomethyl)-1,3-mercaptopropanol.

Example 10

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3,3-Bis(N-ethylperfluorooctanesulfonamidomethyl)oxetane (described in EP-A-0 430 887) is treated with thiourea and sulfuric acid in tert-butanol and subsequently cleaved with 2-(2-aminoethylamino)ethanol as described in Example 1 to yield 2,2-bis(N-ethylperfluorooctanesulfonamidomethyl)-1,3-mercaptopropanol.

Examples 11 to 19

Using the general methods described in Examples 1-10, the following additional mercapto-alcohols are prepared.

Ex.	Fluorochemical	Mercapto-Alcohol
11	CF ₃ CF ₂ CH ₂ SH	(CF ₃ CF ₂ CH ₂ SCH ₂) ₂ C(CH ₂ OH)(CH ₂ SH)
12	C ₆ F ₁₃ (CH ₂) ₄ SH	(C ₆ F ₁₃ (CH ₂) ₄ SO ₂ CH ₂) ₂ C(CH ₂ OH)(CH ₂ SH)
13	C ₈ F ₁₇ CH ₂ CH ₂ CH ₂ SH	(C ₈ F ₁₇ CH ₂ CH ₂ CH ₂ SCH ₂) ₂ C(CH ₂ OH)(CH ₂ SH)
14	C ₈ F ₁₇ CH ₂ CH ₂ N(CH ₃)CH ₂ CH ₂ CH ₂ SH	(C ₈ F ₁₇ CH ₂ CH ₂ N(CH ₃)CH ₂ CH ₂ CH ₂ SCH ₂) ₂ C(CH ₂ OH)(CH ₂ SH)
15	C ₈ F ₁₇ SO ₂ NHCH ₂ CH ₂ SH	(C ₈ F ₁₇ SO ₂ NHCH ₂ CH ₂ SCH ₂) ₂ C(CH ₂ OH)(CH ₂ SH)
16	C ₈ F ₁₇ CH ₂ CH ₂ SO ₂ NHCH ₂ CH ₂ SH	(C ₈ F ₁₇ CH ₂ CH ₂ SO ₂ NHCH ₂ CH ₂ SCH ₂) ₂ C(CH ₂ OH)(CH ₂ SH)
17	C ₇ F ₁₅ CONHCH ₂ CH ₂ S	(C ₇ F ₁₅ CONHCH ₂ CH ₂ SCH ₂) ₂ C(CH ₂ OH)(CH ₂ SH)
18	C ₈ F ₁₇ CH ₂ CH ₂ OH	(C ₈ F ₁₇ CH ₂ CH ₂ OCH ₂) ₂ C(CH ₂ OH)(CH ₂ SH)
19	C ₆ F ₁₃ CH ₂ CH ₂ OH	(C ₆ F ₁₃ CH ₂ CH ₂ OCH ₂) ₂ C(CH ₂ OH)(CH ₂ SH)

Examples 20-21

Stability of Bis-perfluoroalkylmercapto-alcohol Derivatives

Thermogravimetric analyses are run on the mercapto-alcohol of Example 7 and the maleic acid derivative of Example 6. The instrument is run at 10°C/minute to either 350°C or 500°C with 100 ml/min N₂ or in air.

TGA (Wt. Loss)
Temperature (°C) of Indicated Weight Loss

Cmpd of Example	DSC (Tm)	Under Nitrogen			Under Air		
		Init.	10%	50%	Init.	10%	50%
7	165-1	215	286	349	175	255	306
6	28-34	75	261	315	71	250	301

These results indicate the thermal superiority of the subject bis-perfluoroalkylthio-neopentyl type mercapto alcohols and their derivatives. The subject mercapto-alcohol and derivative as exemplified above both the appreciably stable to a temperature greater than 280°C in nitrogen, even to 260°C in air.

Example 22

The mercapto-alcohol of Example 7 is predried azeotropically with isopropyl acetate or with 1,1,1-trichloroethane. An equimolar amount of the dried mercapto-alcohol is then reacted with an equimolar amount of isophorone diisocyanate under nitrogen in the presence of a catalytic amount of triethylamine and dibutyltin dilaurate with isopropyl acetate as solvent. The reaction mixture is heated at reflux for two hours and the completeness of the reaction is indicated by the absence of the N=C=O infrared band at 2270cm⁻¹.

Example 23



2,2-Bis(1,1,2,2-tetrahydroperfluorodecylthiomethyl)-3-bromo-1-propanol (17.4 g, 16.6 mmol; prepared from the corresponding oxetane using hydrobromic acid and described in Example 9 of EP-A-0 430 887), diethyl mercaptosuccinate (6.5 g, 31 mmol), potassium carbonate (4.3 g, 31 mmol) and acetone (50 ml) is charged to a three-necked flask and heated at reflux overnight. Water is added and the mixture is extracted three times with methyl propyl ketone. The product is purified by silica gel chromatography to yield a yellow oil which is 99% pure by GLC.

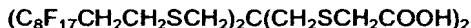
The above diethyl ester (9.5 g, 7.6 mmol) is dissolved in 20 ml of diglyme and water (3 ml) in a single neck flask. Sodium hydroxide (1.5 ml of a 50% aqueous solution) is added and the mixture allowed to stir overnight at room temperature. Water is added, and the mixture extracted with methyl propyl ketone. The aqueous layer is acidified and extracted with methyl propyl ketone. The solvent is removed under vacuum and the resultant yellow oil is precipitated into water to produce a white solid. NMR shows proton resonances at 2.5 ppm (4 x R_fCH₂); 2.6 - 3.0 ppm (10 x SCH₂); 3.51 ppm (2 x CH₂OH); 3.72 ppm (1 x SCHCOO).

Analysis for C₂₉H₂₂F₂₄S₃O₅:

Calculated: C, 29.2; H, 1.8; F, 54.1; S, 8.1.

Found: C, 28.9; H, 1.7; F, 53.9; S, 8.2.

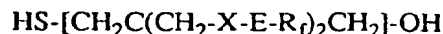
Example 24



2,2-Bis(1,1,2,2-tetrahydroperfluorodecylthiomethyl)-1,3-dibromopropane (25 g, 21 mmol, prepared from the bromopropanol cited in Example 23 using phosphorus tribromide and described in Example 11 of EP-A-0 430 887), ethyl 2-mercaptoacetate (5.1 g, 41 mmol), potassium carbonate (5.8 g, 42 mmol) and acetone (70 ml) is reacted and the product purified as described in Example 23. The resultant diester is saponified, acidified and precipitated into water to yield the above-identified diacid as a white powder.

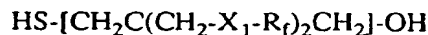
Claims

1. A R_f-neopentyl mercapto-alcohol of formula I or II



(I) or

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(II)

wherein

10 R_f is a straight or branched chain perfluoroalkyl of 1 to 18 carbon atoms or said perfluoroalkyl substituted by perfluoroalkoxy of 2 to 6 carbon atoms,

E is branched or straight chain alkylene of 1 to 10 carbon atoms or said alkylene interrupted by one to three groups selected from the group consisting of -NR-, -O-, -S-, SO_2 -, -COO-, -OOC-, -CONR-, -NRCO-, - SO_2NR -, and - NRSO_2 -, or terminated at the R_f end with -CONR- or - SO_2NR -, where R_f is attached to the carbon or sulfur atom,

15 X is -S-, -O-, - SO_2 -, or -NR-,

X_1 is -CONR- or - SO_2NR -, where R_f is attached to the carbon or sulfur atom, and where

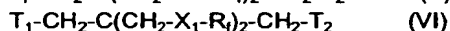
R is independently hydrogen, alkyl of 1 to 6 carbon atoms or hydroxyalkyl of 2 to 6 carbon atoms.

20 2. A compound according to claim 1 where in the compound of formula I or II, R_f is perfluoroalkyl of 4 to 18 carbon atoms or perfluoroalkyl of 2 to 6 carbon atoms substituted by perfluoroalkoxy of 2 to 6 carbon atoms.

3. A compound according to claim 1 where in the compound of formula I, R_f is perfluoroalkyl of 6 to 16 carbon atoms or perfluoroalkyl of 2 to 6 carbon atoms substituted by perfluoroalkoxy of 2 to 6 carbon atoms, E is alkylene of 2 to 6 carbon atoms, -CONHCH₂CH₂-, -CH₂CH₂N(CH₃)CH₂CH₂-, -CH₂CH₂SO₂NHCH₂CH₂-, -CH₂CH₂OCH₂CH₂-, or -SO₂NHCH₂CH₂-, and X is -S-, - SO_2 - or -O-.

30 4. The compound according to claim 1 which is selected from the group consisting of
 $(\text{C}_6\text{F}_{13}\text{CH}_2\text{CH}_2\text{SCH}_2)_2\text{C}(\text{CH}_2\text{OH})(\text{CH}_2\text{SH})$,
 $(\text{C}_8\text{F}_{17}\text{CH}_2\text{CH}_2\text{SCH}_2)_2\text{C}(\text{CH}_2\text{OH})(\text{CH}_2\text{SH})$,
 $(\text{C}_7\text{F}_{15}\text{CONHCH}_2\text{CH}_2\text{SCH}_2)_2\text{C}(\text{CH}_2\text{OH})(\text{CH}_2\text{SH})$,
 $[(\text{CF}_3)_2\text{CFO}(\text{CF}_2)_n\text{CH}_2\text{CH}_2\text{SCH}_2]_2\text{C}(\text{CH}_2\text{OH})(\text{CH}_2\text{SH})$ where n is 3 or 4;
 $[(\text{CF}_3)_2\text{CFOCF}_2\text{CH}_2\text{CH}_2\text{SCH}_2]_2\text{C}(\text{CH}_2\text{OH})(\text{CH}_2\text{SH})$,
 $(\text{C}_8\text{F}_{17}\text{CH}_2\text{CH}_2\text{SO}_2\text{CH}_2)_2\text{C}(\text{CH}_2\text{OH})(\text{CH}_2\text{SH})$,
 $(\text{R}_f\text{CH}_2\text{CH}_2\text{SCH}_2)_2\text{C}(\text{CH}_2\text{OH})(\text{CH}_2\text{SH})$ where
 R_f is a mixture of C_6F_{13} , C_8F_{17} , $\text{C}_{10}\text{F}_{21}$, $\text{C}_{12}\text{F}_{25}$ and $\text{C}_{14}\text{F}_{29}$,
 $(\text{R}_f\text{SO}_2\text{N}(\text{C}_2\text{H}_5)\text{CH}_2)_2\text{C}(\text{CH}_2\text{OH})(\text{CH}_2\text{SH})$ where
 R_f is a mixture of C_6F_{13} , C_8F_{17} , $\text{C}_{10}\text{F}_{21}$, $\text{C}_{12}\text{F}_{25}$ and $\text{C}_{14}\text{F}_{29}$ and
 $(\text{R}_f\text{CH}_2\text{CH}_2\text{OCH}_2)_2\text{C}(\text{CH}_2\text{OH})(\text{CH}_2\text{SH})$ where
 R_f is a mixture of C_6F_{13} , C_8F_{17} , $\text{C}_{10}\text{F}_{21}$, $\text{C}_{12}\text{F}_{25}$ and $\text{C}_{14}\text{F}_{29}$.

45 5. A compound of the formula (V) or (VI)



where

T_1 is the group VII



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where

R_1 , R_2 , R_3 and R_4 are independently hydrogen, alkyl of 1 to 18 carbon atoms, hydroxyalkyl of 1 to 4 carbon atoms, haloalkyl of 1 to 4 carbon atoms, alkoxyalkyl of 2 to 12 carbon atoms, phenyl, phenyl substituted by one or two alkyl of 1 to 4 carbon atoms, by alkoxy of 1 to 4 carbon atoms, by carboxy or by hydroxy, phenylalkyl of 7 to 9 carbon atoms, carboxy, -COOZ where Z is alkyl of 1 to 18 carbon atoms, said alkyl interrupted by one or more -O- groups, allyl, cyclohexyl, phenyl, benzyl, tolyl or naphthyl; or -CONZ₁Z₂ where Z₁ and Z₂ are independently hydrogen, alkyl of 1 to 12 carbon atoms, hydroxyalkyl of 2 to 4 carbon atoms, phenyl, benzyl, alkoxyalkyl of 2 to 12 carbon atoms, cyclohexyl or Z₁ and Z₂ together are pen-

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tamethylene or 3-oxapentamethylene, with the proviso that at least one of R_1 to R_4 contains a carboxy, -COOZ or -CONZ₁Z₂ group,

T_2 has the same meaning as T_1 or is hydroxyl and

R_f is a straight or branched chain perfluoroalkyl of 1 to 18 carbon atoms or said perfluoroalkyl substituted by perfluoroalkoxy of 2 to 6 carbon atoms,

E is branched or straight chain alkylene of 1 to 10 carbon atoms or said alkylene interrupted by one to three groups selected from the group consisting of -NR-, -O-, -S-, SO_2 -, -COO-, -OOC-, -CONR-, -NRCO-, - SO_2NR -, and - $NRSO_2$ -, or terminated at the R_f end with -CONR- or - SO_2NR -, where R_f is attached to the carbon or sulfur atom,

X is -S-, -O-, - SO_2 -, or -NR-,

X_1 is -CONR- or - SO_2NR -, where R_f is attached to the carbon or sulfur atom, and where

R is independently hydrogen, alkyl of 1 to 6 carbon atoms or hydroxyalkyl of 2 to 6 carbon atoms.

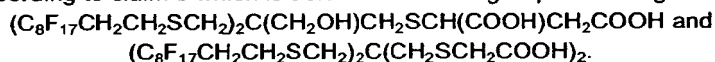
6. A compound according to claim 5 of formula V where

R_f is perfluoroalkyl of 6 to 16 carbon atoms or perfluoroalkyl of 2 to 6 carbon atoms substituted by perfluoroalkoxy of 2 to 6 carbon atoms,

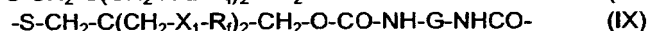
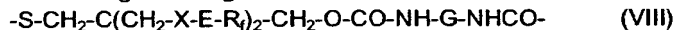
E is alkylene of 2 to 6 carbon atoms, -CONHCH₂CH₂-, -CH₂CH₂N(CH₃)CH₂CH₂-, -CH₂CH₂SO₂NHCH₂CH₂-, -CH₂CH₂OCH₂CH₂-, or -SO₂NHCH₂CH₂-, and

X is -S-, SO_2 - or -O-.

7. The compound according to claim 5 which is selected from the group consisting of



8. A linear polythiourethane having recurring structural units of formula VIII or IX



where

R_f is a straight or branched chain perfluoroalkyl of 1 to 18 carbon atoms or said perfluoroalkyl substituted by perfluoroalkoxy of 2 to 6 carbon atoms,

E is branched or straight chain alkylene of 1 to 10 carbon atoms or said alkylene interrupted by one to three groups selected from the group consisting of -NR-, -O-, -S-, SO_2 -, -COO-, -OOC-, -CONR-, -NRCO-, - SO_2NR -, and - $NRSO_2$ -, or terminated at the R_f end with -CONR- or - SO_2NR -, where R_f is attached to the carbon or sulfur atom,

X is -S-, -O-, - SO_2 -, or -NR-,

X_1 is -CONR- or - SO_2NR -, where R_f is attached to the carbon or sulfur atom, and where

R is independently hydrogen, alkyl of 1 to 6 carbon atoms or hydroxyalkyl of 2 to 6 carbon atoms and

G is the radical of an aliphatic, cycloaliphatic or aromatic diisocyanate after removal of the two NCO groups.

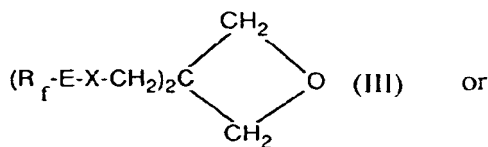
9. A polythiourethane according to claim 8 where in the compound of formula VIII,

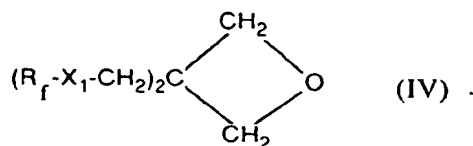
R_f is perfluoroalkyl of 6 to 16 carbon atoms or perfluoroalkyl of 2 to 6 carbon atoms substituted by perfluoroalkoxy of 2 to 6 carbon atoms,

E is alkylene of 2 to 6 carbon atoms, -CONHCH₂CH₂-, -CH₂CH₂N(CH₃)CH₂CH₂-, -CH₂CH₂SO₂NHCH₂CH₂-, -CH₂CH₂OCH₂CH₂-, or -SO₂NHCH₂CH₂-, and

X is -S-, - SO_2 - or -O-.

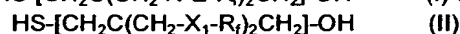
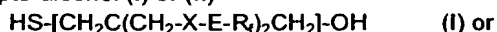
10. A process for the preparation of a compound of formula (I) or (II) of one of claims 1 to 4 which process is characterized by the reaction of a sulfur containing reagent with a bis-perfluoroalkyl oxetane of formula III or IV



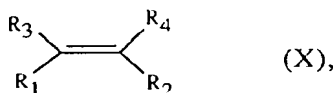


11. A process for the preparation of a compound of formula (V) or (VI) where T_2 is hydroxyl and T_1 is a group VII of one of claims 5 to 7 which process is characterized by

a) the addition of a mercapto-alcohol (I) or (II)

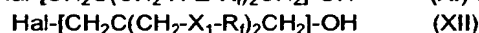
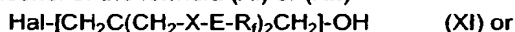


wherein X, X_1 , E and, R_f are defined as hereinbefore,
with a compound of formula (X)



wherein R_1 , R_2 , R_3 and R_4 are defined as hereinbefore or

b) by the reaction of an alcohol of the formula (XI) or (XII)

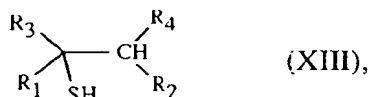


wherein

X, X_1 , E and, R_f are defined as hereinbefore and

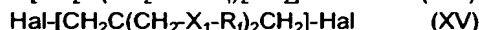
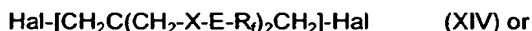
Hal is halogen, preferably chlorine or bromine,

and a thiol of formula (XIII)



wherein R_1 , R_2 , R_3 and R_4 are defined as hereinbefore.

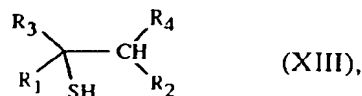
12. A process for the preparation of a compound of formula (V) or (VI) where T_2 has the same meaning as T_1 , according to one of claims 5 to 7 which process is characterized by the addition of a bis-halogen compound (XIV) or (XV)



wherein

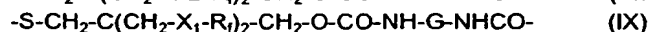
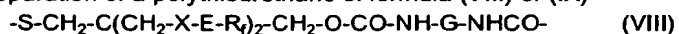
X, X_1 , E and, R_f are defined as hereinbefore, and

Hal is halogen, preferably chloro, bromo or iodo, to a compound of formula and a thiol of formula (XIII)

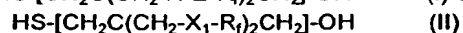
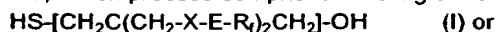


wherein R_1 , R_2 , R_3 and R_4 are defined as hereinbefore.

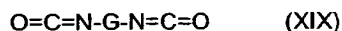
13. A process for the preparation of a polythiourethane of formula (VIII) or (IX)



according to one of claims 8 or 9, which process comprises reacting a mercapto-alcohol of formula I or II



wherein R_f , X, E and X_1 are defined as in one of claims 1 to 4 with an aliphatic, cycloaliphatic or aromatic diisocyanate of formula XIX



wherein

5 G is a bivalent aliphatic, cycloaliphatic or aromatic radical.

14. A method of improving oil and water repellency to substrates which method comprises treating the substrate with a compound of one of claims 1 to 9.
- 10 15. Leather, paper, wood, masonry, metals, plastics, glass, painted surfaces, textiles or metallic surfaces that are treated with an oil and water repellent amount of a compound of one of claims 1 to 9.
16. A method of improving soil repellency to substrates or compositions which method comprises treating the substrate with a polythiourethane of one of claims 8 to 9 or adding a polythiourethane of one of claims 8 or 9 to the composition.
- 15 17. A composition with improved soil repellency which composition comprises a polythiourethane of one of claims 8 or 9.
18. A composition with improved levelling, wetting and spreading properties, which composition comprises a polythiourethane of one of claims 8 or 9.
- 20 19. Formulations for cleaning glass and other hard, non-porous materials, hair care products, paint, stain and varnish formulations, comprising polythiourethane of one of claims 8 or 9.
- 25 20. Papers, wallpaper, asphalt laminates, liner board, cardboard and papers derived from synthetic fibers with improved abrasive properties, comprising a polythiourethane of one of claims 8 or 9.

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European Patent
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EUROPEAN SEARCH REPORT

Application Number

EP 92 81 0118

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl.5)
D,A	US-A-3 886 201 (R. A. FALK ET AL) ---		C07C323/12
A	EP-A-0 348 350 (CIBA-GEIGY) ---		C07C323/25
A	EP-A-0 376 882 (CIBA-GEIGY) ---		C07C323/41
P,A	EP-A-0 430 887 (CIBA-GEIGY) ---		C07C323/49
P,A	EP-A-0 453 406 (CIBA-GEIGY) -----		C07C323/52
			C07C323/65
			C08G75/04
			D06M15/19
			TECHNICAL FIELDS SEARCHED (Int. Cl.5)
			C07C
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 14 APRIL 1992	Examiner ZAROKOSTAS K.
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